

**PCT**WORLD INTELLECTUAL PROPERTY ORGANIZATION  
International Bureau

## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<b>(51) International Patent Classification <sup>7</sup> :</b> <b>G03F 7/00, B29C 33/60</b>	<b>A1</b>	<b>(11) International Publication Number:</b> <b>WO 00/00868</b> <b>(43) International Publication Date:</b> 6 January 2000 (06.01.00)
<b>(21) International Application Number:</b> PCT/US99/14859 <b>(22) International Filing Date:</b> 30 June 1999 (30.06.99)  <b>(30) Priority Data:</b> 09/107,006 30 June 1998 (30.06.98) US  <b>(71) Applicant:</b> REGENTS OF THE UNIVERSITY OF MINNESOTA [US/US]; Morrill Hall, 100 Church Street S.E., Minneapolis, MN 55455 (US). <b>(72) Inventor:</b> CHOU, Stephen, Y.; 7 Foulet Drive, Princeton, NJ 08540 (US). <b>(74) Agent:</b> VIKSNINS, Ann, S.; Schwegman, Lundberg, Woessner & Kluth, P.O. Box 2938, Minneapolis, MN 55402 (US).		<b>(81) Designated States:</b> CN, JP, European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE).  <b>Published</b> <i>With international search report.</i>
<b>(54) Title:</b> SURFACES WITH RELEASE AGENT  <b>(57) Abstract</b>  Thin coatings (less than and approaching monomolecular coatings) of persistent release materials comprising compounds of the formula: $\text{RELEASE-M}(\text{X})_{n-1}-\text{RELEASE-M}(\text{X})_{n-m-1} \text{Q}_m$ or $\text{RELEASE-M}(\text{OR})_{n-1}-$ , wherein RELEASE is a molecular chain of from 4 to 20 atoms in length, M is a metal atom, semiconductor atom, or semimetal atom; X is halogen or cyano, especially Cl, F, or Br; Q is hydrogen or alkyl group; R is hydrogen, alkyl or phenyl, preferably hydrogen or alkyl of 1 to 4 carbon atoms; and; and n-m 1 is at least 1 particularly for a lithographic method and apparatus for creating ultra-fine (sub-25nm) patterns in a thin film coated on a substrate in which a mold having at least one protruding feature is pressed into a thin film carried on a substrate.		

BEST AVAILABLE COPY

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakhstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia ..	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

## SURFACES WITH RELEASE AGENT

**BACKGROUND OF THE INVENTION****1. Field of the Invention**

The present invention relates to release surfaces, particularly release surfaces with fine features to be replicated, and to lithography which may be used to produce integrated circuits and microdevices. More specifically, the present invention relates to a process of using an improved mold or microreplication surface that creates patterns with ultra fine features in a thin film carried on a surface of a substrate.

**2. Background of the Art**

10 In many different areas of technology and commercial utility, it is highly desirable that surface be provided with non-stick functionality. The wide range of utility for this type of technology ranges from antistain treatments for fabrics and surfaces (e.g., countertops, stove tops, and the like), to utensils (e.g., cooking or laboratory utensils and surfaces), release surfaces for imaging technology (e.g., image transfer surfaces, temporary carriers), and mold release surfaces. Antistick functionality has clear lubricating implications where the antistick function can be provided in a substantive or retentive manner onto a substrate.

In the fabrication of semiconductor integrated electrical circuits, integrated optical, magnetic, mechanical circuits and microdevices, and the like, one of the key processing methods is lithography and especially photolithography. Lithography can be used, along with its traditional resist imaging in the formation of printing plates and resist images, to create a pattern in a thin film carried on a substrate so that, in subsequent process steps, the pattern can be replicated in the substrate or in another material which is added onto the substrate. The thin film which accepts a pattern or image during the lithographic process is often referred to as resist. The resist may be either a positive resist or a negative resist, depending on its operation of formation. For example, a positive photoresist becomes more soluble in a solvent where irradiated and a negative resist becomes more insoluble where irradiated. A typical lithographic process for integrated circuit fabrication involves exposing or irradiating a photoresist composition or film with a beam of radiation or

particles, including light, energetic particles (which may be electrons), photons, or ions, by either passing a flood beam through a mask or scanning a focused beam. The radiation or particle beam changes the chemical structure of the exposed area of the film, so that when washed or immersed in a developer or washed with a developer, either the exposed area or the unexposed area of the resist will be removed to recreate the patterns or its obverse of the mask or the scanning. The lithography resolution is limited by the wavelength of the particles and the resolution of the beam, the particle scattering in the resist and the substrate, and the properties of the resist.

There is an ongoing need in art of lithography to produce progressively smaller pattern sizes while maintaining cost efficiency in the process. There is a great need to develop low-cost technologies for mass producing sub-50 nm structures since such a technology could have an enormous impact in many areas of engineering and science. Not only will the future of semiconductor integrated circuits be affected, but also the commercialization of many innovative electrical, optical, magnetic, mechanical microdevices that are far superior to current devices will rely on the possibility of such technology. Additionally optical materials, including reflective coatings and reflective sheeting (as may be used for security purposes or for signage) can use microreplication techniques according to lithographic technology.

Numerous technologies have been developed to service these needs, but they all suffer serious drawbacks and none of them can mass produce sub-50 nm lithography at a low cost. Electron beam lithography has demonstrated 10 nm lithography resolution. A.N. Broers, J.M. Harper, and W.W. Molzen, *Appl. Phys. Lett.* 33, 392 (1978) and P.B. Fischer and S.Y. Chou, *Appl. Phys. Lett.* 62, 2989 (1993). However, using these technologies for mass production of sub-50 nm structures seems economically impractical due to inherent low throughput in a serial processing tool. X-ray lithography, which can have a high throughput, has demonstrated 50 nm lithography resolution. K. Early, M.L. Schattenburg, and H.I. Smith, *Microelectronic Engineering* 11, 317 (1990). But X-ray lithography tools are rather expensive and its ability for mass producing sub-50 nm structures is yet to be commercially demonstrated. Lithography based on scanning probes has produced sub-10 nm structures in a very thin layer of

materials. However, the practicality of such lithography as a manufacturing tool is hard to judge at this point.

Imprint technology using compressive molding of thermoplastic polymers is a low cost mass manufacturing technology and has been around for several decades. Features with sizes greater than 1 micrometers have been routinely imprinted in plastics. Compact disks which are based on imprinting of polycarbonate are one example of the commercial use of this technology. Other examples are imprinted polymethyl methacrylate (PMMA) structures with a feature size on the order to 10 micrometers for making micromechanical parts.

10 M. Harmening, W. Bacher, P. Bley, A. El-Kholi, H. Kalb, B. Kowanz, W. Menz, A. Michel, and J. Mohr, *Proceedings IEEE Micro Electro Mechanical Systems*, 202 (1992). Molded polyester micromechanical parts with feature dimensions of several tens of microns have also been used. H. Li and S.D. Senturia, *Proceedings of 1992 13th IEEE/CHMT International Electronic Manufacturing*

15 *Technology Symposium*, 145 (1992). However, no one has recognized the use of imprint technology to provide 25 nm structures with high aspect ratios. Furthermore, the possibility of developing a lithographic method that combines imprint technology and other technologies to replace the conventional lithography used in semiconductor integrated circuit manufacturing has never

20 been raised.

#### **SUMMARY OF THE INVENTION**

The present invention relates to methods for changing the properties of surfaces by bonding coatings of molecules to surfaces to form non-continuous coatings of molecules bonded thereto. The invention is particularly

25 advantageous for forming mold or microreplication surfaces having coatings of molecules bonded thereto, and to processes of molding and microreplication using these coatings and surfaces. The coatings may be referred to as non-continuous coatings as the coating material does not have to bond cohesively with itself parallel to the surface which is coated, but is bonded, molecule-by-

30 molecule, to the surface, such as grass protrudes, blade-by-blade, from the surface of the ground.

The present invention relates to a method for providing a surface with a treatment that can render the surface more effective in molding or

microreplication processes. A molecular moiety having release properties towards other materials (e.g., fluorinated hydrocarbon chains or polysiloxanes) and low chemical reactivity to moldable polymers is bonded to a mold or microreplication surface. The release properties of the molecular moiety having

5 release properties allows for the enhancement of resolution on the molded article since the molded material is released from the minute features of the mold on a molecular level. More common polymeric coated release surfaces can fill the openings or partially fill the openings of the mold. Merely smoother release surfaces expose the surface of the mold to abrasion and to reaction with the

10 molding materials. The description of the coating as non-continuous may be described as follows. A continuous coating normally is one that forms a film on the surface with no direct route from one side of the film to the other side of the film. As there is no true film coating formed in the practice of the present invention, but rather individual molecules tend to be stacked up on the surface,

15 there is no continuous coating, even though there may be uniform properties over the surface. On a molecular level, the surface would appear as a surface having one moiety at one end of a relatively linear molecule bonded to the surface. The relatively linear molecule extends away from the surface, with the release properties provided by the 'tail' of the molecule that extends away from

20 the surface. The relative concentration of tails on the surface controls the hydrophilic/hydrophobic/polar/non-polar properties of the surface so that it will enable ready release of the material provided by the molding or microreplication process. The release portion of the adhered molecule will preferably have few reactive sites on the tail, particularly within the last one, two, three or four

25 skeletal atoms in the relatively linear chain (e.g., with a hydrocarbon-based chain, the alpha, beta, gamma, and delta atoms in the chain). Such moieties to be avoided particularly would include free hydrogen containing groups (e.g., acid groups, carboxylic acid groups or salts, sulfonic acid groups or salts, amine groups, ethylenically unsaturated groups, and the like).

30 The present invention also relates to a method and apparatus for performing ultra-fine line lithography of the type used to produce integrated circuits and microdevices. A layer of thin film is deposited upon a surface of a substrate. A mold having its mold surface treated with the release materials of

the present invention and at least one protruding feature and a recess is pressed into the thin film, therefore the thickness of the film under the protruding feature is thinner than the thickness of the film under the recess and a relief is formed in the thin film. The relief generally conforms to the shape of the feature on the mold. After the mold is removed from the film, the thin film is processed such that the thinner portion of the film in the relief is removed exposing the underlying substrate. Thus, the pattern in the mold is replicated in the thin film, completing the lithography. The patterns in the thin film will be, in subsequent processes, reproduced in the substrate or in another material that is added onto the substrate. The use of the release treatment on the mold surface enhances the resolution of the image and can protect the mold so that it can be used more often without showing wear on fine features in the mold.

The invention described here is based on a fundamentally different principle from conventional lithography. The process invention can eliminate many resolution limitations imposed in conventional lithography, such as wavelength limitation, backscattering of particles in the resist and substrate, and optical interference. It has been demonstrated the present invention can include a high throughput mass production lithography method for generating sub-25 nm features. Furthermore, the present invention has the ability to mass produce sub-10 nm features at a low cost. These capabilities of the present invention is unattainable with the prior art, and the use of the adherent release property coating improves the durability and the resolution of the process even further. The present process, however, has implications and utility for more macroscopic details in molding surfaces and would include features in the super-50 nm range, the super-100 nm range, and the super 200 nm range, as well as macroscopic dimensions in the visual range of features (e.g., 0.1 mm and greater).

#### **BRIEF DESCRIPTION OF THE DRAWINGS**

Figure 1A is a cross sectional view showing a mold and substrate in accordance with the present invention.

Figure 1B is a cross sectional view of the mold and substrate of Figure 1A showing the mold pressed into a thin film carried on the substrate.

Figure 1C is a cross sectional view of the substrate of Figure 1B following compression of the mold into the thin film.

Figure 1D is a cross sectional view of the substrate of Figure 1C showing removal of compressed portions of the thin film to expose the underlying substrate.

Figure 2 is a scanning electron micrograph of a top view of 25 nm  
5 diameter holes with a 120 nm period formed by compressive molding into a PMMA film as shown in Figure 1C.

Figure 3 is a scanning electron micrograph of a top view of a 100 nm wide trench formed by compressive molding into a PMMA film as shown in Figure 1C.

10 Figure 4 is a scanning electron micrograph of a perspective view of the strips formed by compressive molding into a PMMA film as shown in Figure 1C. The strips are 70 nm wide and 200 nm tall, and have a high aspect ratio, a surface roughness less than 3 nm, and corners of nearly a perfect 90 degrees.

Figure 5A is a cross sectional view of the substrate of Figure 1D  
15 following deposition of a material.

Figure 5B is a cross sectional view of the substrate of Figure 5A following selective removal of the material by a lift off process.

Figure 6 is a scanning electron micrograph of the substrate of Figure 2 following deposition of material and a lift off process. The metal dots have a 25  
20 nm diameter that is the same as that of the holes created in the PMMA shown in Figure 2.

Figure 7 is a scanning electron micrograph of the substrate of Figure 3 following deposition of material in a lift off process. The metal linewidth is 100 nm that is the same as the width of the PMMA trenches shown in Figure 3.

25 Figure 8 is a cross sectional view of the substrate of Figure 1D following subsequent processing.

Figure 9 is a simplified block diagram of an apparatus in accordance with one embodiment of the invention.

#### **DETAILED DESCRIPTION OF THE INVENTION**

30 The present invention relates to methods for changing the properties of surfaces by bonding non-continuous coatings of molecules thereto, to surfaces having non-continuous coatings of molecules bonded thereto, to mold or microreplication surfaces having non-continuous coatings of molecules bonded



thereto, and to processes of molding and microreplication using these coatings and surfaces.

This invention also relates to a method and apparatus for a high-resolution, high-throughput, low-cost lithography. Unlike current  
5 microlithography, a preferred embodiment of the present invention abandons usage of energetic light or particle beams. Photolithography may also benefit from the practice of the present invention by the use of the reactive release layer bonded to the mold surface. In the embodiment of the invention which does not require the use of photolithography, the present invention is based on pressing a  
10 mold into a thin film on a substrate to create a relief and, later removing the compressed area of the film to expose the underlying substrate and to form a resist pattern on the substrate that replicates the obverse of the protruding pattern of the mold.

The present invention also has demonstrated the generation of patterns,  
15 such as holes, pillars, or trenches in a thin film on a substrate, that have a minimum size of 25 nm, a depth over 100 nm, a side wall smoothness better than 3 nm, and corners with near perfect 90 degrees angles. It was found that presently the size of imprinted features is limited by the size of the mold being used; with a suitable mold, the present invention should create sub-10 nm  
20 structures with a high aspect ratio. Furthermore, using one embodiment of the present invention that including a material deposition and a lift-off process, 100 nm wide metal lines of a 200 nm period and 25 nm diameter metal dots of 125 nm period have been fabricated. The resist pattern created using the present invention also has been used as a mask to etch nanostructures (features having  
25 dimensions less than 1000 nm, preferably less than 500 nm) into the substrate.

The present invention offers many unique advantages over the prior art. First, since it is based on a paradigm different from the prior art and it abandons the usage of an energetic particle beam such as photons, electrons, and ions, the present invention eliminates many factors that limit the resolution of  
30 conventional lithographies, such as wave diffraction limits due to a finite wavelength, the limits due to scattering of particles in the resist and the substrate, and interferences. Therefore the present invention offers a finer lithography resolution and much more uniform lithography over entire substrate than the

prior art. Results show it can achieve sub-25 nm resolution. Second, the present invention can produce sub-25 nm features in parallel over a large area, leading to a high throughput. This seems unachievable with the prior art. And thirdly, since no sophisticated energetic particle beam generator is involved, the present invention can achieve a sub-25 nm lithography over a large area at a cost much lower than the prior art. These advantages make the present invention superior to the prior art and vital to future integrated circuit manufacturing and other areas of science and engineering where nanolithography is required.

The non-continuous coatings of molecules are formed from a specific type of reactive compound. These compounds may be characterized by the following structure:



or



RELEASE is a molecular chain of from 4 to 20 atoms in length, preferably from 6 to 16 atoms in length, which molecule has either polar or non-polar properties, depending upon the phobicity desired towards a molding agent;

M is an inorganic atom, especially a metal atom, semiconductor atom, or semimetal atom;

X is halogen or cyano, especially Cl, F, or Br;

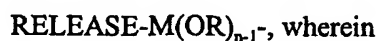
R is hydrogen, alkyl or phenyl, preferably hydrogen or alkyl of 1 to 4 carbon atoms, most preferably hydrogen, methyl or ethyl; and;

(n) is the valence -1 of M, usually 1, 2 or 3 depending upon the nature of M.

The actual moiety bonded to the surface has one of the groups bonded to the metal or semimetal atom removed during a reaction with the mold surface and may have the structural formula:



or



RELEASE is a molecular chain of from 4 to 20 atoms in length, preferably from 6 to 16 atoms in length, which molecule has either polar or non-polar properties;

M is a metal or semimetal atom;

5 X is halogen or cyano, especially Cl, F, or Br;

R is hydrogen, alkyl or phenyl, preferably hydrogen or alkyl of 1 to 4 carbon atoms; and;

(n) is the valence -1 of M.

- 10 As noted above, the properties of RELEASE are determined in part by the nature of the molded material to be used with the surface or the nature of the properties desired on the surface. That is where the surface is to be used in microreplication with a polar polymeric material, the RELEASE properties must be non-polar. Non-polar RELEASE groups are preferably selected, for example,
- 15 from non-polar molecular units including especially siloxane units and highly fluorinated or fluorocarbon units. It is further preferred that these non-polar molecular units are linear units of from 4 to 20 skeletal atoms in the linear chain. Smaller chains might not form as continuous of release properties as desired, and longer chains might mask features on the surface to be replicated. By highly
- 20 fluorinated is meant that at least 2/3 of all substituents on the carbon are fluorinated units, with the remaining units comprising Cl or H. Preferably the terminal carbon is perfluorinated, more preferably the terminal carbon atom is perfluorinated and no hydrogen atoms are present on the three terminal carbon atoms, and most preferably the chain is perfluorinated.
- 25 M is preferably a metal atom, semiconductor atom or semimetal atom such as for example, Si, Ti, Zr, Cr, Ge, and the like. Most preferably M is Si. In these cases, n would preferably be 3.

Examples of the compounds which can be used in the practice of the present invention comprise perfluorohexyl trichlorosilane, perfluorooctyl

30 trichlorosilane, perfluorodecyl trichlorosilane, perfluorododecyl trichlorosilane, perfluorohexylpropyl trichlorosilane, perfluorodecyl trichlorotitanium, perfluorodecyl dichlorobromosilane, polydimethylsiloxane-trichlorosilane (with n preferably of about 4 to 20 for the polydimethylsiloxane unit), perfluorodecyl

dichlorobromogermanium, perfluorodecyl dichlorobromochromium, and the like.

The mold surfaces to be used may be any surface to which the release providing molecules may bond. By selecting appropriate release providing molecules, substantially any release surface may be used. The release surface may be metallic, semimetallic, metal oxides, metal and semimetal carbides and nitrides, semimetallic oxide, polymeric, semiconductors, photocinductors, ceramic, glass, composite or the like, as is known in the molding and microreplication art. Particularly useful substrates include, but are not limited to, silicon, silicon nitride, silicon carbide, silicon nitride, doped semiconductor blends, photoconductors (both organic and inorganic), and the like. The molding process may include impression molding as generally described above, injection molding, powder molding, blow molding, casting or cast molding, vapor deposition molding, decomposition molding (where materials are decomposed to form new materials which deposit on the surface), and the like. Uniformly shaped patterns or random patterns may be manufactured, and the materials used in the molding composition may harden, as previously noted, by cooling thermally softened materials, polymerizable materials, chemically reacting materials, vapor depositing materials, or the like. Preferred materials comprise semiconductor, dielectric, photoresponsive, thermally responsive, or electrically responsive substrates or surfaces, such as, but not limited to, inorganic oxides (or sulfides, halides, carbides, nitrides, etc.), rare earth oxides (or sulfides, halides, carbides, nitrides, etc.), inorganic or organic silicon compounds (e.g., silica oxides, sulfides, halides, carbides, nitrides, etc.) and their titanium, germanium, cadmium, zinc and the like counterparts (e.g., titania, zinc oxide [particles or layers], germanium oxide, cadmium sulfide) as continuous or discontinuous coatings or layers, as mixture, dispersions or blends, as layered structures, and the like.

The release-coating forming materials of the present invention may be applied in coatings which form less than continuous monomolecular layers of the release material. That is, the release material forms coatings comprising tails of the release moiety secured to the surface by reaction with the nominatively inorganic end of the molecule (e.g., the silicon, titanium, germanium, end). The

entire surface of the substrate is not necessarily coated, as the release molecules tend to prevent other molecules from aligning uniformly (at least uniformly in a pattern) along the surface. There may be, and most likely always is, some spacing between the individual coating molecules on the surface since, as shown in Figure 1A, the coating does not form as a continuous layer parallel to the coated surface, but rather forms as extended molecules bonded at only one end to the surface, leaving the RELEASE group outwardly extending to provide the release (non-stick) properties. However, the release moiety tail of the compounds evidences an area of lubricity, so a uniform coating is not essential.

Coating weights of the release coating material may be used in surprisingly small amounts, considering their effectiveness. For example, coating weights of less than  $0.001\text{mg/m}^2$  of surface area have provided significant release coating effects. Coating weights of 0.001 to 100 or more  $\text{mg/m}^2$  of surface area, from 0.005 to  $5\text{mg/m}^2$  of surface area, and preferably from 0.01 up to 1 to  $5\text{mg/m}^2$  of surface area are generally useful.

Figures 1A-1D show steps in accordance with one embodiment. Figure 1A shows molding layer 10 having body 12 and molding layer 14. The release coating material Si-RELEASE is shown attached to said molding layer 10, although not proportionally. The Si-RELEASE compound is shown as single molecules bonded at the Si end, with the RELEASE tail extending therefrom to provide the release properties to the mold 14. The size of the release compound residues -Si-RELEASE is molecular as opposed to the macromolecular view of the molding surface 14 shown in the Figure 1A. The residual groups which may be attached to the Si (e.g., unreacted H, cyano, or halogen) are not shown, merely for convenience in drawing the Figure. As can be seen from this less than literal representation, the RELEASE moieties extend away from the molding surface 14. These RELEASE 'tails' provide the release property and tend to be fairly durable and persistent. Molding layer 14 is shown as including a plurality of features 16 having a desired shape. A release layer 17 is shown bonded to the surface of the features 16 on the molding layer 14. A substrate 18 carries thin film layer 20. Thin film layer 20 is deposited through any appropriate technique such as spin casting, slot die coating, slide coating, curtain coating, solvent

coating, gravure coating, screen coating, vapor deposition, sputtering and the like.

Figure 1B shows a compressive molding step where mold 10 is pressed into thin film layer 20 in the direction shown by arrow 22 forming compressed regions 24. In the embodiment, shown in Figures 1A-1D, features 16 are not pressed all of the way into thin film 20 and do not contact substrate 18. In some embodiments, top portions 24a of film 20 may contact depressed surfaces 16a of mold 10. This causes top surfaces 24a to substantially conform to the shape of surfaces 16a, for example flat. When contact occurs, this also can stop the mold move further into the thin film 20, due to a sudden increase of contact area and hence a decrease of the compressive pressure when the compressive force is constant. The release layer 17 of the present inventions improves the release of the thin film layer 20 from the features 16 of the mold 10.

Figure 1C is a cross sectional view showing thin film layer 20 following removal of mold 10. Layer 20 includes a plurality of recesses formed at compressed regions 24 which generally conform to the shape of features 16 which is coated with release layer 17. Layer 20 is subjected to a subsequent processing step as shown in Figure 1D, in which the compressed portions 24 of film 20 are removed thereby exposing substrate 18. This removal may be through any appropriate process such as reactive ion etching, wet chemical etching. This forms dams 26 having recesses 28 on the surface of substrate 18. Recesses 28 form relief features that conform generally to the shape of features 16 and mold 10.

The mold 10 is patterned with features 16 comprising pillars, holes and trenches with a minimum lateral feature size of 25 nm, using electron beam lithography, reactive ion etching (RIE) and other appropriate methods. The typical depth of feature 16 is from 5 nm to 200 nm (either including the dimensions of the release layer 17 or excluding those molecular dimensions), depending upon the desired lateral dimension. In general, the mold should be selected to be hard relative to the softened thin film, and can be made of metals, dielectrics, polymers, or semiconductors or ceramics or their combination. In one experiment, the mold 10 consists of a layer 14 and features 16 of silicon dioxide on a silicon substrate 12.

Thin film layer 20 may comprise a thermoplastic polymer or other thermoplastic, hardenable, or curable material which may pass from a flowable state to a non-flowing state upon a change in conditions (e.g., temperature, polymerization, curing or irradiation). During the compressive molding step shown in Fig. 1B, thin film 20 may be heated at a temperature to allow sufficient softening of the film relative to the mold. For example, above the glass transition temperature the polymer has a low viscosity and can flow, thereby conforming to the features 16 without forming a strong adherence to the surface because of the presence of the release layer 17. The film layer may comprise anything from continuous films of materials, to lightly sintered materials, to loose powders held in place by gravity until the compressive and adherent steps of the molding or microreplication processes. For example, the material could be a polymer film, latex film, viscous polymer coating, composite coating, fusible powder coating, blend of adherent and powder, lightly sintered powder, and the like. The polymer may comprise any moldable polymer, including, but not limited to (meth)acrylates (which includes acrylates and methacrylates), polycarbonates, polyvinyl resins, polyamides, polyimides, polyurethanes, polysiloxanes, polyesters (e.g., polyethyleneterephthalate, polyethylenenaphthalate), polyethers, and the like. Materials such as silica, alumina, zirconia, chromia, titania, and other metal oxides (or halides) or semimetal oxides (or halides) whether in dry form or sol form (aqueous, inorganic solvent or organic solvent) may be used as the moldable material. Composites, mixing both polymeric materials and non-polymeric materials, including microfibers and particulates, may also be used as the molding material.

In one experiment, the thin film 20 was a PMMA spun on a silicon wafer 18. The thickness of the PMMA was chosen from 50 nm to 250 nm. PMMA was chosen for several reasons. First, even though PMMA does not adhere well to the SiO<sub>2</sub> mold due to its hydrophilic surface, its adherence can be reduced further by the use of the release layers of the present invention. Good mold release properties are essential for fabricating nanoscale features. Second, shrinkage of PMMA is less than 0.5% for large changes of temperature and pressure. See I. Rubin, Injection Molding, (Wiley, New York) 1992. In a molding process, both the mold 10 and PMMA 20 were first heated to a

temperature of 200°C which is higher than the glass transition temperature of PMMA, 105°C. See M. Harmening, W. Bacher, P. Bley, A. El-Kholi, H. Kalb, B. Kowanz, W. Menz, A. Michel, and J. Mohr, Proceedings IEEE Micro Electro Mechanical Systems, 202 (1992). Then the mold 10 and features 16 were  
5 compressed against the thin film 20 and held there until the temperature dropped below the PMMA's glass transition temperature. Various pressures have been tested. It was found that the one preferred pressure is about 400-1900 psi., especially 500-100 psi. At that pressure, the pattern of the features 16 can be fully transferred into the PMMA, particularly when the release was expedited by  
10 the presence of the release layer 17. After removing mold 10, the PMMA in the compressed area was removed using an oxygen plasma, exposing the underlying silicon substrate and replicating the patterns of the mold over the entire thickness of the PMMA. The molding pressure is, of course, dependent upon the specific polymer being used and can therefore vary widely from material to material.

15 Figure 2 shows a scanning electron micrograph of a top view of 25 nm diameter holes with a 120 nm period formed into a PMMA film in accordance with Figure 1C. Mold features as large as tens of microns on the same mold as the nanoscale mold features have been imprinted.

Figure 3 shows a scanning electron micrograph of a top view of 100 nm  
20 wide trenches with a 200 nm period formed in PMMA in accordance with Figure 1C.

Figure 4 is a scanning electron micrograph of a perspective view of trenches made in the PMMA using the present invention with embodiment that top portions 24a of film 20 contact depressed surfaces 16a of mold 10. The  
25 strips are 70 nm wide, 200 nm tall, therefore a high aspect ratio. The surface of these PMMA features is extremely smooth and the roughness is less than 3 nm. The corners of the strips are nearly a perfect 90 degrees. Such smoothness, such sharp right angles, and such high aspect ratio at the 70 nm features size cannot be obtained with the prior art.

30 Furthermore, scanning electron microscopy of the PMMA patterns and the mold showed that the lateral feature size and the smoothness to the sidewalls of PMMA patterns fabricated using the present invention conform with the mold. From our observations, it is clear that the feature size achieved so far with the



present invention is limited by our mold size. From the texture of the imprinted PMMA, it appears that 10 nm features can be fabricated with the present invention.

After the steps 1A-1D, the patterns in film 20 can be replicated in a material that is added on substrate 18 or can be replicated directly into substrate 18. Figures 5A and 5B show one example of the subsequent steps which follow the steps of Figures 1A-1D. Following formation of the recesses 28 shown in Figure 1D, a layer of material 30 is deposited over substrate 18 as shown in Figure 5A. Material 30 is deposited through any desired technique over dams 26 and into recesses 28 between dams 26. Material 30 may comprise, for example, electrical conductors or semiconductors or dielectrics of the type used to fabricate integrated circuits, or it may comprise ferromagnetic materials for magnetic devices. Next, a lift off process is performed in which a selective chemical etch is applied which removes dams 26 causing material 30 deposited on top of dams 26 to be removed. Figure 5B shows the structure which results following the lift off process. A plurality of elements 32 formed of material 30 are left on the surface of substrate 18. Elements 32 are of the type used to form miniaturized devices such as integrated circuits. Subsequent processing steps similar to those shown in steps 1A-1D may be repeated to form additional layers on substrate 18.

Figure 6 is a scanning electron micrograph of the substrate of Figure 2 following deposition of 5 nm of titanium and 15 nm of gold and a lift off process. In the lift-off process, the wafers were soaked in acetone to dissolve the PMMA and therefore lift-off metals which were on the PMMA. The metal dots have a 25 nm diameter that is the same as that of the holes created in the PMMA using the present invention.

Figure 7 is a scanning electron micrograph of the substrate of Figure 3 following deposition of 5 nm of titanium and 15 nm of gold and a lift off process. The metal linewidth is 100 nm that is the same as the width of the PMMA trenches shown in Figure 3. Figures 6 and 7 have demonstrated that, during the oxygen RIE process in the present invention, the compressed PMMA area was completely removed and the lateral size of the PMMA features has not been changed significantly.

Figure 8 is a cross sectional view of substrate 18 of Figure 1D following an example alternative processing step that replicates the patterns in film 20 directly into substrate 18. In Figure 8, substrate 18 has been exposed to an etching process such as reactive ion etching, chemical etching, etc., such that  
5 recesses 40 are formed in substrate 18. These recesses 40 may be used for subsequent processing steps. For example, recesses 40 may be filled with material for use in fabricating a device. This is just one example of a subsequent processing step which can be used in conjunction with the present invention.

Molding processes typically use two plates to form malleable material  
10 therebetween. In the present invention, substrate 18 and body 12 (mold 10) act as plates for the imprint process of the invention. Substrate 18 and body 12 should be sufficiently stiff to reduce bending while forming the imprint. Such bending leads to deformation in the pattern formed in the film 20.

Figure 9 is a simplified block diagram of apparatus 50 for performing  
15 nanoimprint lithography in accordance with the invention. Apparatus 50 includes stationary block 52 carrying substrate 18 and moveable molding block 54 carrying mold 10. Blocks 52 and 54 carry the substrate 18 and mold 10 depicted in Figures 1A-1D. A controller 56 couples to x-y positioner 58 and z positioner 60. An alignment mark 64 is on mold 10 and complimentary mark 68  
20 is on substrate 18. Sensor 62 carried in block 54 couples to alignment marks 64 and 68 and provide an alignment signal to controller 56. Controller 56 is also provided with input output circuitry 66.

In operation, controller 56 controls the imprinting of mold 10 into film 20 on substrate 18 by actuating z positioner 60 which moves block 54 in the z  
25 direction relative to block 52. During the imprinting process, precise alignment of mold 10 and film 20 is crucial. This is achieved using optical or electrical alignment techniques. For example, sensor 62 and alignment marks 64 and 68 may be an optical detector and optical alignment marks which generate a moiré alignment pattern such that moiré alignment techniques may be employed to  
30 position mold 10 relative to film 20. Such techniques are described by Nomura et al. A MOIRÉ ALIGNMENT TECHNIQUE FOR MIX AND MATCH LITHOGRAPHIC SYSTEM, J. Vac. Sci. Technol. B6(1), Jan/Feb 1988, pg. 394 and by Hara et al., AN ALIGNMENT TECHNIQUE USING DEFRACATED

MOIRÉ SIGNALS J. Vac. Sci. Technol. B7(6), Nov/Dec 1989, pg. 1977.

Controller 56 processes this alignment information and adjusts the position of block 54 in the x-y plane relative to film 20 using x-y positioner 58. In another embodiment, alignment marks 64 and 68 comprise plates of a capacitor such that  
5 sensor 62 detects capacitance between marks 64 and 68. Using this technique, alignment is achieved by moving block 54 in the x-y plane to maximize the capacitance between alignment marks 64 and 68. During imprinting, controller 56 may also monitor and control the temperature of film 20.

It should be understood that the invention is not limited to the specific  
10 technique described herein, and may be implemented in any appropriate lithographic process. Generally, the mold should be hard relative to the film during the molding process. This may be achieved for example, by sufficiently heating the film. Additionally, it should be understood that the invention is not limited to the particular film described herein. For example, other types of films  
15 may be used. In one alternative embodiment, a thin film may be developed which has a chemical composition which changes under pressure. Thus, following the imprint process, a chemical etch could be applied to the film which selectively etches those portions whose composition had changed due to applied pressure. In another embodiment, after molding of the thin film to create a  
20 thickness contrast in the thin film, a material is deposited on the thin film and the thickness contrast then is transferred into the substrate.

Although the present invention has been described with reference to preferred embodiments, workers skilled in the art will recognize that changes may be made in form and detail without departing from the spirit and scope of  
25 the invention.

### **EXAMPLES**

An example of a lithographic process according to the present invention forming a pattern in a film carried on a substrate would be practiced by the steps of depositing a film on a substrate to provide a mold having a protruding feature  
30 and a recess formed thereby, the feature and the recess having a shape forming a mold pattern. At least a portion of the surface, (in this case a surface of silica or silicon-nitride is preferred) such as the protruding feature(s), if not the entire surface (the protrusions and valleys between the protrusions) onto which the film

is deposited, is coated with the release material comprises a material having the formula:

RELEASE-M(X)<sub>n-1</sub>-, Formula I

RELEASE-M(X)<sub>n-m-1</sub>Q<sub>m</sub> Formula II

5 or

RELEASE-M(OR)<sub>n-1</sub>-, Formula III wherein

RELEASE is a molecular chain of from 4 to 20 atoms in length, preferably from 6 to 16 atoms in length, which molecule has either polar or non-  
10 polar properties;

M is a metal or semimetal atom;

X is halogen or cyano, especially Cl, F, or Br;

Q is a hydrogen or alkyl group,

m is the number of Q groups,

15 R is hydrogen, alkyl or phenyl, preferably hydrogen or alkyl of 1 to 4 carbon atoms; and;

n-m-1 in Formula II is at least 1 (m is 2 or less), preferably 2 (m is 1 or less), and most preferably at least 3 (m is 0)

n is the valence -1 of M.

20 In particular, silicon compounds (pure or in solution) of C 1 to C4 alkyl (for R), wherein X is F, and RELEASE is perfluorinated alkyl are preferred. Particularly 1H, 1H, 2H, 2H-perfluorododecyltrichlorosilane (commercially available as a 97% solids solution) has been found to be particularly useful in the practice of the invention. (The triethoxysilane counterpart tends to require a more active  
25 stimulus to assure extensive bonding to the surface. The 1H, 1H, 2H, 2H-perfluorododecylmethyldichlorosilane, would close in effectiveness to the 1H, 1H, 2H, 2H-perfluorododecyltrichlorosilane, with the slightly reduced activity of the additional methyl group replacing one of the chloro groups on the silane. Similarly, the commercially available 1H, 1H, 2H, 2H-  
30 perfluorododecyltrimethylmonochlorosilane would be slightly less reactive, yet again). This 1H, 1H, 2H, 2H-perfluorododecyltrichlorosilane compound is coated (in a room temperature, air tight, ventilated environment) at about 0.01mg/m<sup>2</sup> of surface area, heated (to about 40 to 50 degrees Centigrade) to react

the material to the surface, and cooled. This forms a coating on the surface in which the reactive portion of the molecule (the SiF bonds) reacts with the silica or silica nitride surface, forming a coating comprising the silicon atom bonded to the surface with a tail of the perfluorinatedalkyl group extending from the surface to leave a reduced friction surface. The mold is then urged into the film whereby the thickness of the film under the protruding feature is reduced and a thin region is formed in the film. The mold is removed from the film, processing the relief. The thin region is removed, exposing a portion of the surface of the substrate which underlies the thin region. The exposed portion of the surface of the substrate substantially replicates the mold pattern. The improvement of having at least a portion of said protruding feature and a portion of said release having the release materials of the invention bonded thereto improves the release and the resolution of the mold operation. Importantly, the release coating of the invention has been proven to be persistent and reusable, particularly where modest pressures (e.g., less than 1000 psi are used, and where the film does not contain ingredients which chemically attack the release coating. The selection of the release coating with perfluorinated R groups assists in providing chemical attack resistant coatings. It is important to note that the processes and release coated materials of the present invention can be made by the simple coating and reaction of the release coating forming materials of the present invention, and that these materials, and the broad range of equivalents are broadly enabled. The materials may be coated as pure material and allowed to react at ambient conditions (where the materials are particularly active to the surface), they may be in solution to dilute the coating (taking care to select solvents which are themselves not active to the release-coating forming compounds and preferably not to the surface), their reaction may be accelerated by heat, catalysts, initiators (either thermal, or photoinitiators, for example, such as fluorinated sulfonic acids, sulfonium or iodonium photoinitiators with complex halide anions, such as triarylsulfonium hexafluoroantimonate, diaryl iodonium tetrafluoroborate), accelerators and the like.

The release-forming coatings of the present invention may be applied as release coatings by simply applying the chemical compounds to a surface to which they react (essentially any surface with free Hydrogen atoms, which react

with halogens, organic acids, silicic or inorganic acids, hydroxyl groups, hydrogen-containing amine groups, mercaptan groups, and the like). The surfaces may be polymeric surfaces, metallic surfaces, alloy surfaces, ceramic surfaces, composite surfaces, organic surfaces, inorganic surfaces, smooth surfaces, rough surfaces, textured surfaces, patterned surfaces, and the like. The use of temperatures and solvents is limited solely by their effect on the substrate and the coating. That is temperatures should not be used during the application of the surface which would degrade the surface or the coating material or so rapidly volatilize the coating material that it would not adhere. As noted elsewhere, catalysts and initiators may be used, but the preferred release coating forming compounds of the invention generally can react at room temperature without any significant stimulus being applied.

The 1H, 1H, 2H, 2H-perfluorododecyltrichlorosilane has been applied as a release surface to Si surfaces, SiN surfaces and the like solely by application of the commercially available 1H, 1H, 2H, 2H-perfluorododecyltrichlorosilane (without modification) to the surface at room temperature. The compounds of Formula I are the most preferred (primarily because of their activity), the compounds of Formula II less preferred, and the compounds of Formula III least preferred because of their reduced reactivity to surfaces.

WHAT IS CLAIMED IS:

1. A lithographic method for forming a pattern in a film carried on a substrate, comprising the steps of:
- 5 depositing a film on a substrate to provide a mold having a protruding feature and a recess formed thereby, the feature and the recess having a shape forming a mold pattern;
- urging the mold into the film whereby the thickness of the film under the protruding feature is reduced and a thin region is formed in the film;
- 10 removing the mold from the film;
- processing the relief whereby the thin region is removed exposing a portion of the surface of the substrate which underlies the thin region; and
- whereby the exposed portion of the surface of the substrate substantially replicates the mold pattern,
- 15 the improvement comprising at least a portion of said protruding feature and a portion of said release have bonded thereto a release material comprising an inorganic linking group bonded to a molecular chain having release properties.
2. The method of claim 1 wherein said release material comprises a material
- 20 having the formula:
- $$\text{RELEASE-M(X)}_{n-1}-$$
- or
- $$\text{RELEASE-M(OR)}_{n-1}-, \text{ wherein}$$
- 25 RELEASE is a molecular chain of from 4 to 20 atoms in length, preferably from 6 to 16 atoms in length, which molecule has either polar or non-polar properties;
- M is a metal or semimetal atom;
- X is halogen or cyano, especially Cl, F, or Br;
- 30 R is hydrogen, alkyl or phenyl, preferably hydrogen or alkyl of 1 to 4 carbon atoms; and;
- (n) is the valence -1 of M.

3. The process of claim 2 wherein RELEASE comprises a highly fluorinated organic group.
4. The process of claim 3 wherein said highly fluorinated group comprises a  
5 perfluorinated alkyl group.
5. The process of claim 2 wherein X is halogen.
6. The process of claim 2 wherein X is chloro or bromo.
- 10 7. The process of claims 1, 2, 3, 4, 5 or 6 wherein M is Si.
8. A process for improving the release properties of a surface comprising:
  - (a) providing a surface;
  - 15 (b) contacting that surface with a release forming material wherein said release forming material comprises a material having the formula:
 
$$\text{RELEASE-M(X)}_{n-1}-$$

or

$$\text{RELEASE-M(OR)}_{n-1}-, \text{ wherein}$$
- 20 RELEASE is a molecular chain of from 4 to 20 atoms in length, preferably from 6 to 16 atoms in length, which molecule has either polar or non-polar properties;
 

M is a metal or semimetal atom;

X is halogen or cyano, especially Cl, F, or Br;

- 25 R is hydrogen, alkyl or phenyl, preferably hydrogen or alkyl of 1 to 4 carbon atoms; and;
 

(n) is the valence  $-1$  of M.
- 9. The process of claim 8 wherein said release forming material comprises a  
30 material of the formula:
 
$$\text{RELEASE-M(X)}_{n-1}-$$

wherein



RELEASE is a molecular chain of from 4 to 20 atoms in length, which molecular chain has either polar or non-polar properties;

M is a metal or semimetal atom;

X is halogen or cyano, especially Cl, F, or Br; and

5 (n) is the valence -1 of M.

10. The process of claim 9 wherein RELEASE comprises a highly fluorinated organic group.

10 11. The process of claim 10 wherein said highly fluorinated group comprises a perfluorinated organic group.

12. The process of claim 11 wherein said highly fluorinated group comprises a perfluorinated alkyl group of from 4 to 16 carbon atoms.

15

13. The process of claims 8, 9, 10 11 or 12 wherein M is Si.

14. A surface having good anti-adherent properties comprising a surface having bonded thereto the residue of material of the formula:

20 RELEASE-M(X)<sub>n-1</sub>-

or

RELEASE-M(OR)<sub>n-1</sub>-, wherein

RELEASE is a molecular chain of from 4 to 20 atoms in length, preferably from 6 to 16 atoms in length, which molecule has either polar or non-

25 polar properties;

M is a metal or semimetal atom;

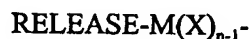
X is halogen or cyano, especially Cl, F, or Br;

R is hydrogen, alkyl or phenyl, preferably hydrogen or alkyl of 1 to 4 carbon atoms; and;

30 (n) is the valence -1 of M,

said material being bonded to said surface only through a bond directly to M where a group X or OR has been removed to enable bonding to said surface, with the group RELEASE still attached to M.

15. The surface of claim 14 wherein said material comprises a material of the formula:



5 wherein

RELEASE is a molecular chain of from 4 to 20 atoms in length, preferably from 6 to 16 atoms in length, which molecule has either polar or non-polar properties;

M is a metal or semimetal atom;

10 X is halogen or cyano, especially Cl, F, or Br; and

(n) is the valence -1 of M.

16. The surface of claim 15 wherein RELEASE comprises a highly fluorinated group.

15

17. The surface of claim 16 wherein said highly fluorinated group comprises a perfluorinated group.

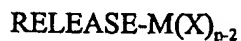
18. The surface of claim 17 wherein said highly fluorinated group comprises a perfluorinated group of from 4 to 16 carbon atoms.

20

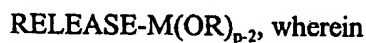
19. The surface of claims 14, 15, 16, 17 or 18 wherein M is Si.

20. A surface having good antiadherent properties comprising a surface having bonded thereto the material of the formula:

25



or



RELEASE is a molecular chain of from 4 to 20 atoms in length, preferably from 6 to 16 atoms in length, which molecule has either polar or non-polar properties;

30

M is a metal or semimetal atom;

X is halogen or cyano, especially Cl, F, or Br;

R is hydrogen, alkyl or phenyl, preferably hydrogen or alkyl of 1 to 4 carbon atoms; and;

p is the valence of M,

said material being bonded to said surface only through a bond directly to M.

5

21. The surface of claim 20 wherein the surface comprises a patterned mold surface.

22. The method of claim 1 wherein the feature on the mold is formed from material selected from the group consisting of: semiconductors, dielectrics, metals, ceramics, polymers and their combination.

23. The surface of claim 14 wherein said surface comprises a material selected from the group consisting of: semiconductors, dielectrics, metals, ceramics, polymers and their combination.

24. The method of claim 1 wherein the step of processing comprises reactive ion etching.

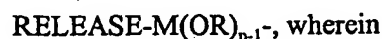
25. The method of claim 1 including repeating the steps of obtaining a mold, urging, removing, and processing to form a multilayered device.

26. A lithographic method for forming a pattern in a film carried on a substrate, comprising the steps of:

obtaining a substrate having a release coating thereon formed by the reaction of a compound of the formula



or



RELEASE is a molecular chain of from 4 to 20 atoms in length, preferably from 6 to 16 atoms in length, which molecule has either polar or non-polar properties;

M is a metal or semimetal atom;

X is halogen or cyano, especially Cl, F, or Br;

R is hydrogen, alkyl or phenyl, preferably hydrogen or alkyl of 1 to 4 carbon atoms; and;

(n) is the valence -1 of M,

5 depositing a film on the substrate;

obtaining a mold having a protruding feature and a recess formed thereby, the feature and the recess having a shape forming a mold pattern; urging the mold into the film creating a thickness contrast pattern in the film;

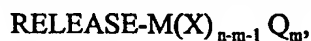
10 removing the mold from the film; and

transferring the thickness contrast pattern in the film onto the substrate.

27. A process for improving the release properties of a surface comprising:

(a) providing a surface;

15 (b) contacting that surface with a release forming material wherein said release forming material comprises a material having the formula:



RELEASE is a molecular chain of from 4 to 20 atoms in length, preferably from 6 to 16 atoms in length, which molecule has either polar or non-  
20 polar properties;

M is a metal or semimetal atom;

X is halogen or cyano, especially Cl, F, or Br;

Q is a hydrogen or alkyl group,

m is the number of Q groups,

25 n-m-1 is at least 1, and

n is the valence -1 of M.

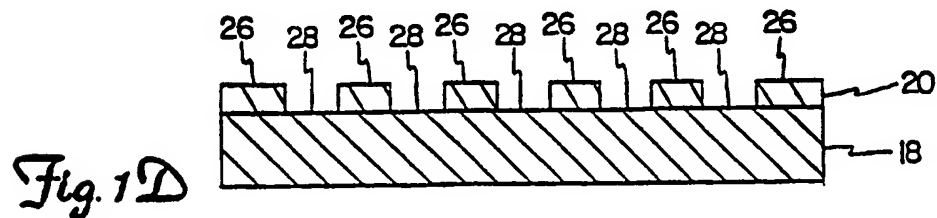
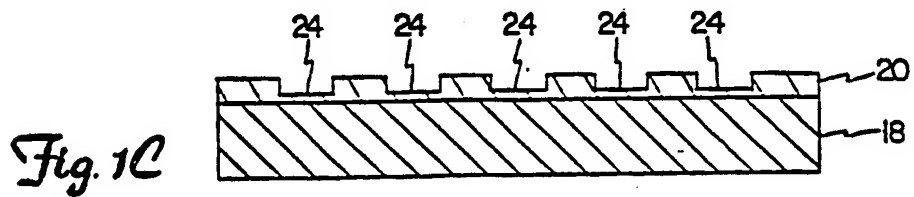
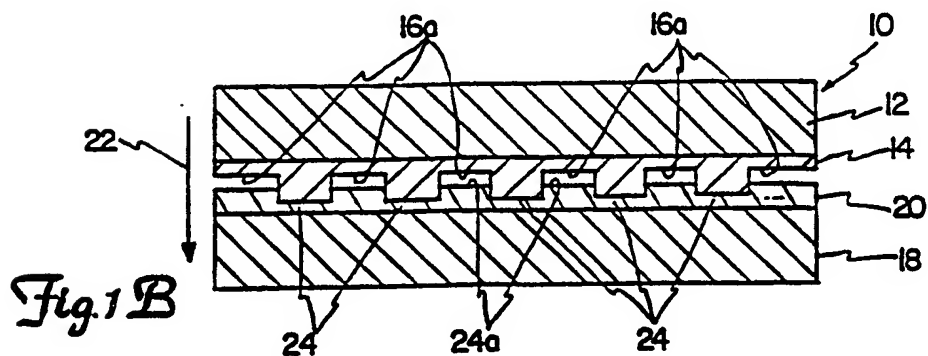
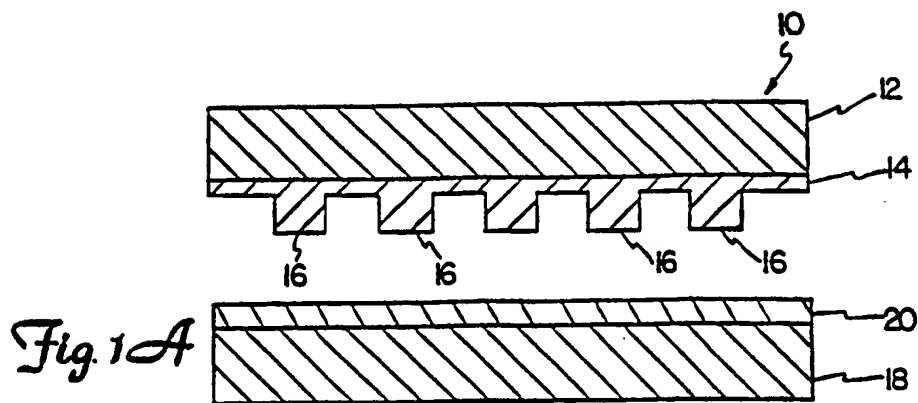
28. The process of claim 27 wherein said release material comprises a material having the formula:

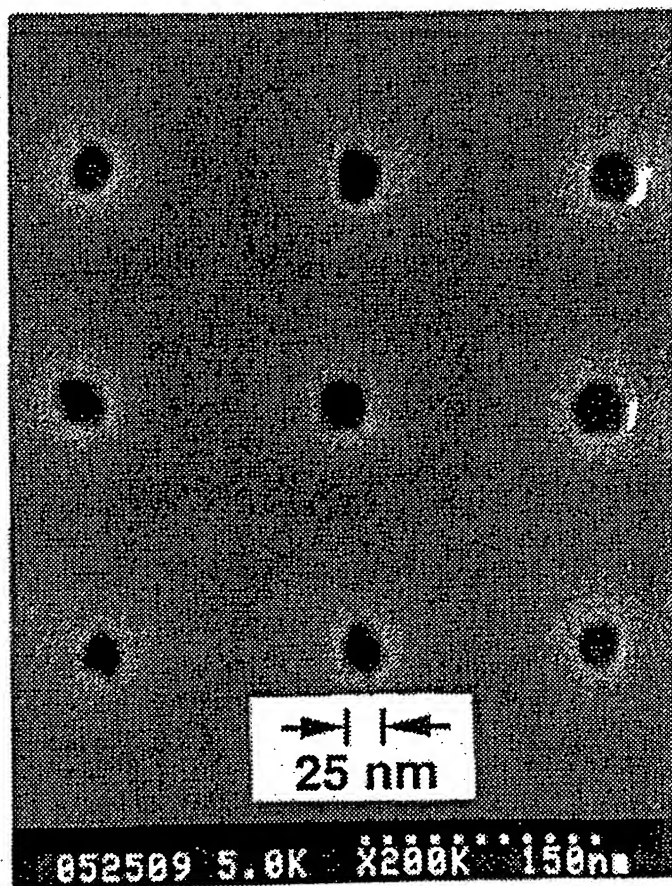


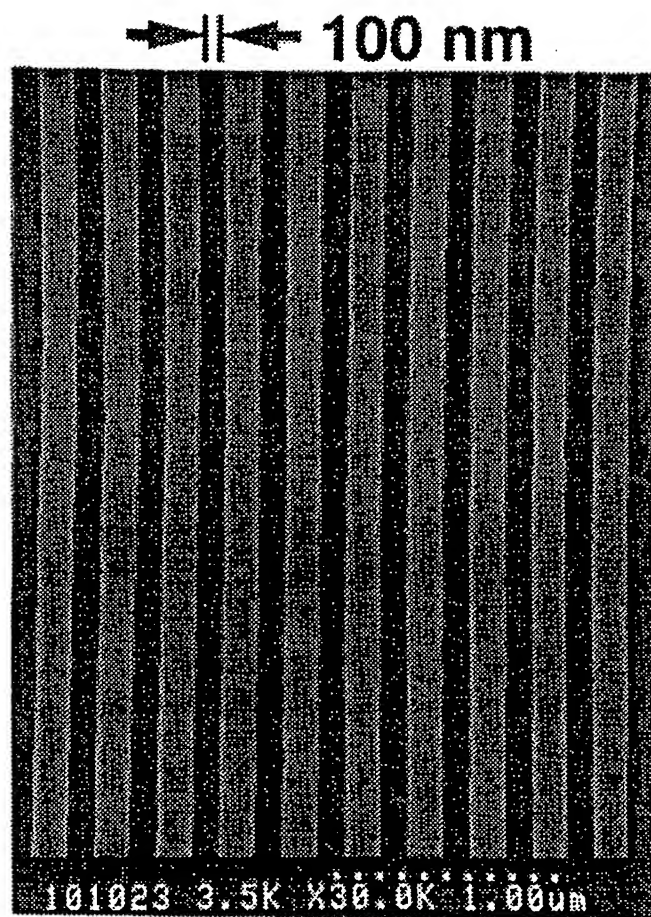
29. The process of claim 38 wherein M is Si;

X is halogen Cl or Br;

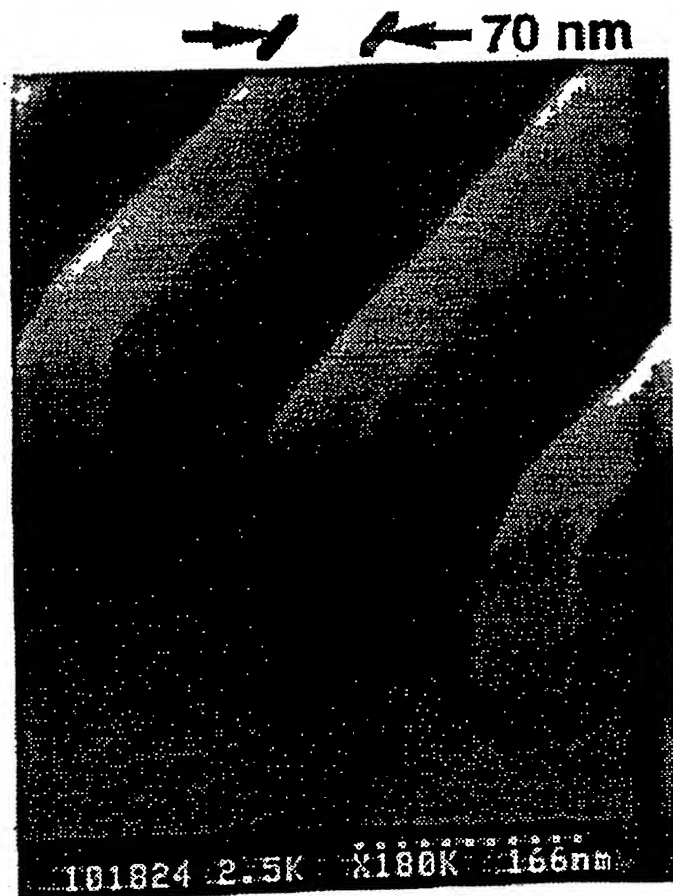
RELEASE is perfluoroalkyl of 6 to 20 carbon atoms;  
and  
n is 3.



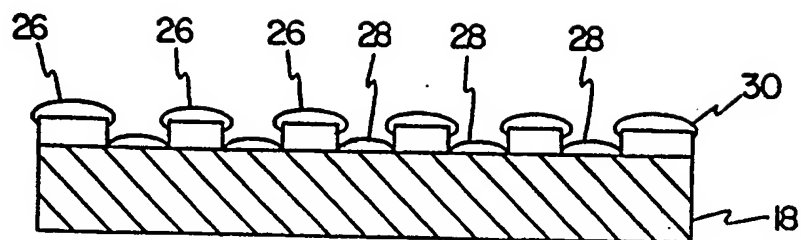
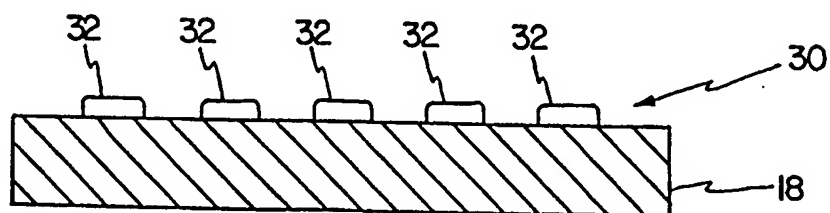
*Fig. 2*

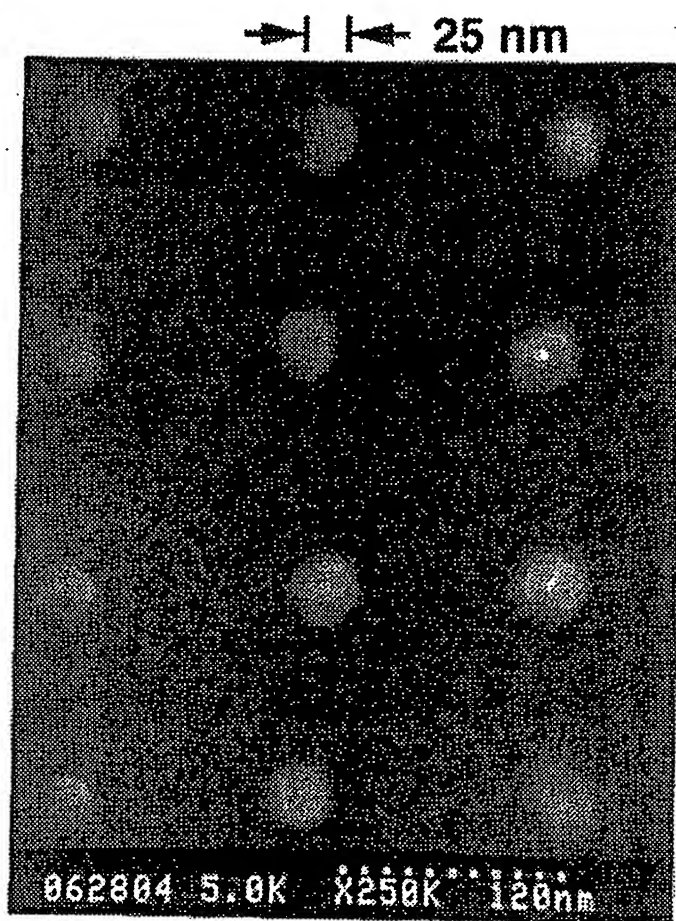
*Fig. 3*

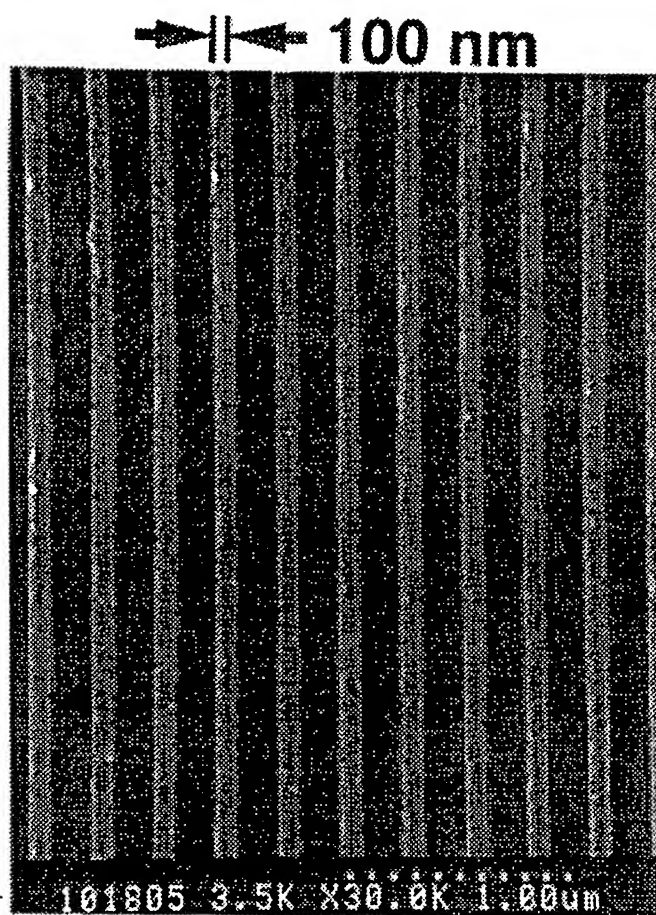




*Fig. 4*

*Fig. 5A**Fig. 5B*

*Fig. 6*

*Fig. 7*

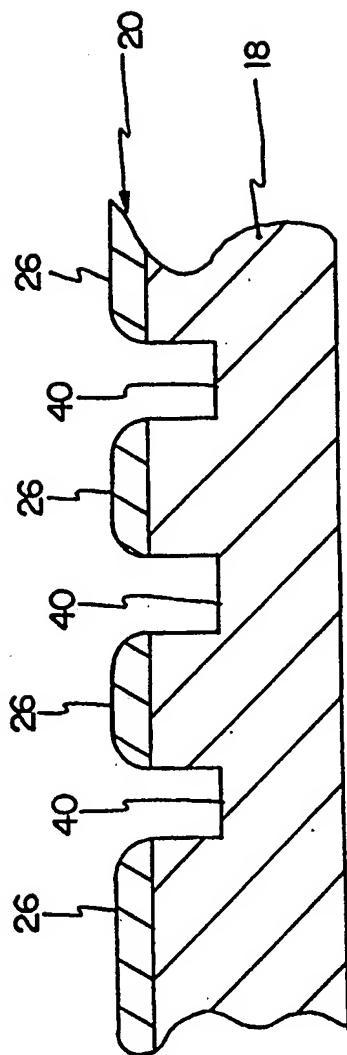
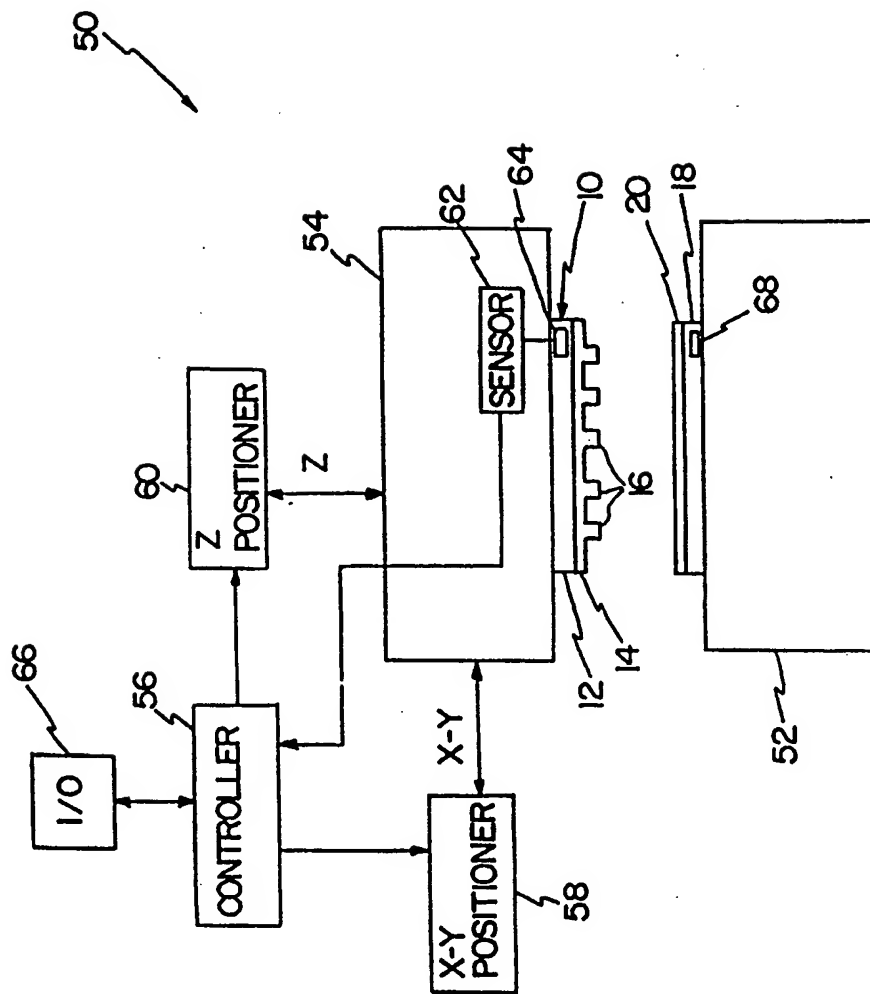


Fig. 8

*Fig. 9*

# INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/14859

**A. CLASSIFICATION OF SUBJECT MATTER**  
IPC 7 G03F7/00 B29C33/60

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)  
IPC 7 G03F B29C

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	EP 0 244 884 A (PHILIPS NV) 11 November 1987 (1987-11-11)	1-7, 23-26
X	the whole document	8-21, 27-29
Y	US 4 731 155 A (NAPOLI LOUIS S ET AL) 15 March 1988 (1988-03-15) the whole document * specially Col.5, L.3-25 *	1-7, 23-26
X	PATENT ABSTRACTS OF JAPAN vol. 017, no. 034 (M-1357), 22 January 1993 (1993-01-22) & JP 04 255307 A (MATSUSHITA ELECTRIC IND CO LTD), 10 September 1992 (1992-09-10) abstract	8-21, 27-29
	-/-	

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

**\* Special categories of cited documents :**

- \*A\* document defining the general state of the art which is not considered to be of particular relevance
- \*E\* earlier document but published on or after the international filing date
- \*L\* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- \*O\* document referring to an oral disclosure, use, exhibition or other means
- \*P\* document published prior to the international filing date but later than the priority date claimed

- \*T\* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- \*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- \*Y\* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- \*&\* document member of the same patent family

Date of the actual completion of the international search

7 October 1999

Date of mailing of the international search report

18/10/1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  
Fax: (+31-70) 340-3016

Authorized officer

Haenisch, U

# INTERNATIONAL SEARCH REPORT

International Application No  
PCT/US 99/14859

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	PATENT ABSTRACTS OF JAPAN vol. 013, no. 495 (P-956), 9 November 1989 (1989-11-09) & JP 01 196749 A (HOYA CORP), 8 August 1989 (1989-08-08) abstract ---	
A	PATENT ABSTRACTS OF JAPAN vol. 017, no. 177 (M-1393), 6 April 1993 (1993-04-06) & JP 04 332694 A (MATSUSHITA ELECTRIC IND CO LTD), 19 November 1992 (1992-11-19) abstract ---	
A	WO 98 26913 A (ERICSSON TELEFON AB L M) 25 June 1998 (1998-06-25) -----	



# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 99/14859

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 0244884	A	11-11-1987	NL 8600809 A	16-10-1987
			JP 7096229 B	18-10-1995
			JP 62236713 A	16-10-1987
			US 4775554 A	04-10-1988
US 4731155	A	15-03-1988	NONE	
JP 04255307	A	10-09-1992	JP 2106048 C	06-11-1996
			JP 8018336 B	28-02-1996
JP 01196749	A	08-08-1989	NONE	
JP 04332694	A	19-11-1992	NONE	
WO 9826913	A	25-06-1998	SE 508968 C	23-11-1998
			AU 5505398 A	15-07-1998
			SE 9604675 A	20-06-1998

THIS PAGE BLANK (USPTO)

**This Page is Inserted by IFW Indexing and Scanning  
Operations and is not part of the Official Record**

**BEST AVAILABLE IMAGES**

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

☐ BLACK BORDERS

☐ IMAGE CUT OFF AT TOP, BOTTOM OR SIDES

☐ FADED TEXT OR DRAWING

☒ BLURRED OR ILLEGIBLE TEXT OR DRAWING

☐ SKEWED/SLANTED IMAGES

☒ COLOR OR BLACK AND WHITE PHOTOGRAPHS

☐ GRAY SCALE DOCUMENTS

☐ LINES OR MARKS ON ORIGINAL DOCUMENT

☐ REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY

☐ OTHER: \_\_\_\_\_

**IMAGES ARE BEST AVAILABLE COPY.**

**As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.**

THIS PAGE BLANK (USPTO)